

# Scaling of dynamic spin correlations in $\text{BaCu}_2(\text{Si}_{0.5}\text{Ge}_{0.5})_2\text{O}_7$

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The magnetic dynamic structure factor of the one-dimensional  $S = 1/2$  chain system  $\text{BaCu}_2(\text{Si}_{0.5}\text{Ge}_{0.5})_2\text{O}_7$  is studied in a wide range of energy transfers and temperatures. Contrary to previous erroneous reports [T. Masuda *et al.*, Phys. Rev. Lett. **93**, 077206 (2004)], the scaling properties observed in the range 0.5–25 meV are found to be fully consistent with expectations for a Luttinger spin liquid. At higher energies, a breakdown of scaling laws is observed and attributed to lattice effects. The results are complementary to those found in literature for other  $S = 1/2$ -chain compounds, such as  $\text{KCuF}_3$  and Cu-benzoate.

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## I. INTRODUCTION

The two isostructural oxides  $\text{BaCu}_2\text{Ge}_2\text{O}_7$  and  $\text{BaCu}_2\text{Si}_2\text{O}_7$  are prototypical quasi-one dimensional (quasi-1D) quantum  $S = 1/2$  antiferromagnets (AF).<sup>1</sup> Previous extensive neutron scattering experiments on  $\text{BaCu}_2\text{Si}_2\text{O}_7$  were instrumental in developing an understanding of the spin dynamics in weakly-coupled  $S = 1/2$  chains.<sup>2,3,4,5</sup> More recently, solid solutions of type  $\text{BaCu}_2(\text{Si}_{1-x}\text{Ge}_x)_2\text{O}_7$  were recognized as potential spin chain systems with random bond strengths.<sup>6</sup> Bulk measurements<sup>6,7,8</sup> and preliminary neutron scattering studies<sup>6</sup> indicated that the  $x = 0.5$  compound shows some very unusual scaling of the static magnetic susceptibility and the dynamic spin correlation function. It was proposed that, due to intrinsic quenched structural disorder,  $\text{BaCu}_2(\text{Si}_{0.5}\text{Ge}_{0.5})_2\text{O}_7$  is an experimental realization of the much-studied Random Singlet (RS) model.<sup>9,10</sup> Unfortunately, when followup measurements were performed on larger and higher quality samples, these conclusions were shown to be erroneous.<sup>11</sup> In fact, in a wide energy range the magnetic dynamic structure factor  $S(q, \omega)$  follows theoretical expectations for disorder-free spin chains. In the present paper we report detailed temperature-dependent measurements of magnetic excitations in  $\text{BaCu}_2(\text{Si}_{0.5}\text{Ge}_{0.5})_2\text{O}_7$  obtained using the new large single crystal samples. The focus is on scaling relations for the spin correlation function. Convenient magnitudes of magnetic interactions enable us to cover a wide dynamic range  $\hbar\omega/\kappa T \approx 1 - 100$ . In this, our results are complementary to those previously obtained for two other  $S = 1/2$  chain systems, namely Cu-benzoate<sup>12</sup> and  $\text{KCuF}_3$  (Ref. 13), where  $\hbar\omega/\kappa T \approx 0.1 - 10$ .

$\text{BaCu}_2(\text{Si}_{1-x}\text{Ge}_x)_2\text{O}_7$  crystallizes in an orthorhombic structure, space group  $Pnma$ . The lattice constants linearly depend on Ge content and for  $x = 0.5$  are:  $a = 6.917 \text{ \AA}$ ,  $b = 13.28 \text{ \AA}$  and  $c = 6.944 \text{ \AA}$ . Magnetic properties are due to  $\text{Cu}^{2+}$  ions that form weakly-coupled antiferromagnetic chains running along the crystallographic  $c$ -axis. The in-chain exchange constant is  $J = 24 \text{ meV}$  for the silicate ( $x = 0$ ) and  $J = 50 \text{ meV}$  for the Ge-system ( $x = 1$ ).<sup>1</sup> The average coupling strength  $\langle J \rangle$  for intermediate concentrations can be deduced from the position of the Bonner-Fisher susceptibility maximum. It increases linearly with  $x$ ,<sup>7,8</sup> and for  $\text{BaCu}_2(\text{Si}_{0.5}\text{Ge}_{0.5})_2\text{O}_7$   $\langle J \rangle = 37 \text{ meV}$ . The corresponding Des Cloizeau-Pearson (DCP) zone-boundary energy of magnetic excitations is  $\hbar\omega_{\text{ZB}} = \pi\langle J \rangle/2 = 58 \text{ meV}$ .<sup>14</sup> Nearest-neighbor spin-spin separation within the chains is equal to  $c/2$ , so the 1D AF zone center  $q_{\parallel} = \pi$  corresponds to  $l = 1$ , where  $(h, k, l)$  denotes a vector in crystallographic reciprocal space. Weak interactions between the chains result in long-range AF ordering at  $T_N = 9.2 \text{ K}$  in the silicate and  $T_N = 10 \text{ K}$  in the germanate.<sup>1</sup> However, for intermediate concentrations, the transition temperature is suppressed,<sup>7,8</sup> and for  $\text{BaCu}_2(\text{Si}_{0.5}\text{Ge}_{0.5})_2\text{O}_7$ , is as low as  $T_N = 0.7 \text{ K}$ .<sup>15</sup> Previous theoretical and neutron scattering studies have shown that, in weakly coupled spin chains, 3D effects are totally negligible at energies  $\hbar\omega \gtrsim 5\kappa T_N$ ,<sup>2,16</sup> so that, barring any disorder-induced effects,  $\text{BaCu}_2(\text{Si}_{0.5}\text{Ge}_{0.5})_2\text{O}_7$  can be expected to behave like a good 1D system for  $\hbar\omega \gtrsim 0.5 \text{ meV}$ . Comparing this energy to  $\hbar\omega_{\text{ZB}}$ , we see that the material presents a conveniently broad energy range for testing any predicted 1D scaling relations for  $S(\mathbf{q}, \omega)$  in quantum  $S = 1/2$  chains.

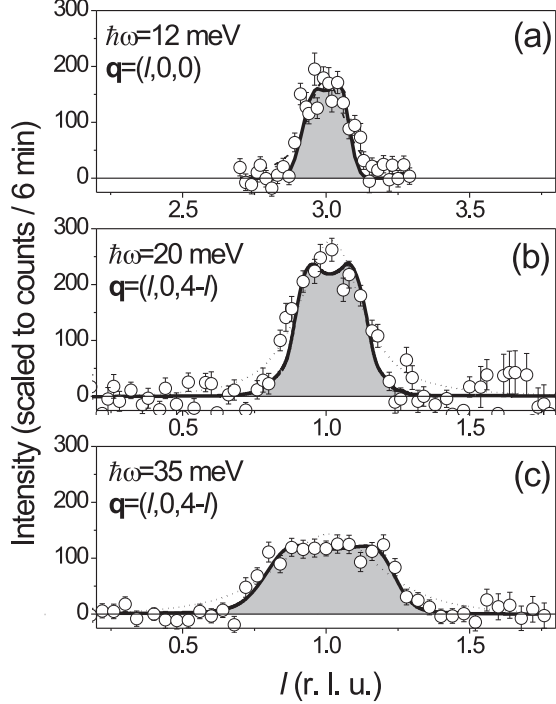


FIG. 1: Constant- $E$  scans measured in  $\text{BaCu}_2(\text{Si}_{0.5}\text{Ge}_{0.5})_2\text{O}_7$  using Setup 1 (3-axis spectrometer) at  $T = 12$  K (symbols). Heavy solid lines and shaded areas are fits using the Müller ansatz. Dashed lines are Lorentzian fits. All fitting functions are convoluted with the known experimental resolution.

## II. EXPERIMENTAL

In the present study we used 6  $\text{BaCu}_2(\text{Si}_{0.5}\text{Ge}_{0.5})_2\text{O}_7$  single crystals of total mass 15 g, grown using the floating zone method. Most crystals were twinned in such a way that the  $a$  and  $c$  axes of the two crystallographic domains were interchanged. Such a growth habit is consistent with  $a \approx c$ , but complicates the measurements, as it involves the chain axis. Individual crystals were co-aligned to form a compound sample with a total effective mosaic of  $1.7^\circ$ . Inelastic neutron scattering data were collected in two separate series of experiments. Experimental Setup 1 employed the HB1 thermal 3-axis spectrometer installed at the High Flux isotope Reactor at ORNL. Final neutron energy was fixed at 14.7 meV. A Pyrolytic Graphite (PG) monochromator was used in conjunction with a PG filter installed after the sample. The collimation setup was  $48' - 40' - 40' - 240'$ . The sample was mounted with the  $(a, c)$  scattering plane horizontal, making crystallographic wave vectors  $(h, 0, l)$  accessible for measurements. Sample environment was a standard He-flow cryostat. All data were corrected for the wavelength-dependent efficiency of the beam monitor (due to higher-order beam contamination) and the

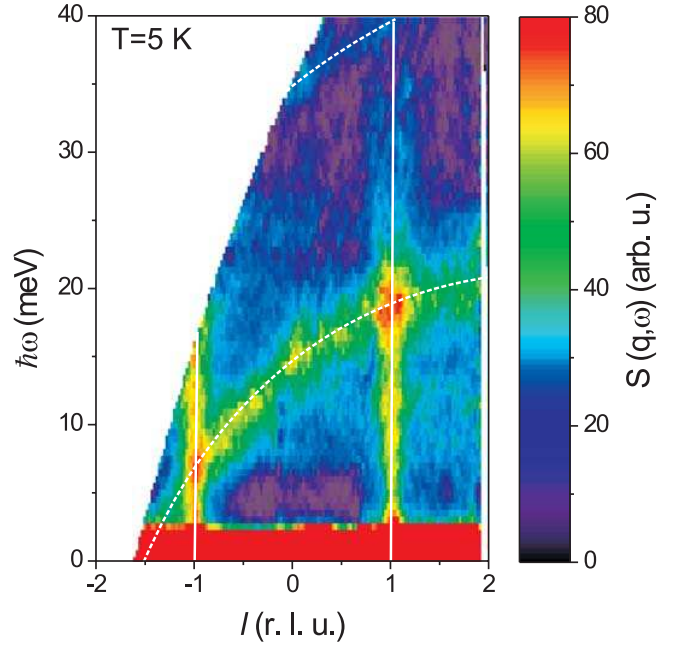


FIG. 2: (Color online) Time of flight spectrum collected in  $\text{BaCu}_2(\text{Si}_{0.5}\text{Ge}_{0.5})_2\text{O}_7$  at  $T = 5$  K using Setup 2, in projection onto the  $c$  (chain) axis. Solid and dashed white lines are 1D antiferromagnetic zone-centers for the two crystallographic domains present in the sample, respectively.

vertical focal length of the fixed-curvature monochromator. It was, in part, an incorrect application of these corrections, that led to erroneous conclusions in Ref. 6. The second series of measurements were performed on the MAPS time-of-flight (TOF) chopper spectrometer installed at the ISIS spallation neutron facility at RAL. Incident energy was fixed at either 50 meV (Setup 2) or 100 meV (Setup 3). The  $(a, c)$  plane of the crystal was mounted parallel to the preferential scattering plane of the instrument. All TOF data were integrated along the  $b^*$  crystallographic direction in the range  $-1 < k < 1$ . Sample temperature was controlled between 5.5 K and 300 K using a closed-cycle refrigerator.

## III. RESULTS AND DATA ANALYSIS

### A. 3-axis data

In 3-axis experiments the data were collected in constant- $E$  scans. Due to the presence of two sets of crystallographic domains, any scattering at wave vector  $(h, k, l)$  is observed simultaneously with that at  $(l, k, h)$ . A steep dispersion of magnetic excitations along the chains implies that, at low energy transfers, all relevant scattering is concentrated near the 1D AF zone-centers  $l$ -odd. For this reason, scans performed at  $l = 0$  effectively pick up the signal from only one of the two domains. While only half of the sample contributes to scattering,

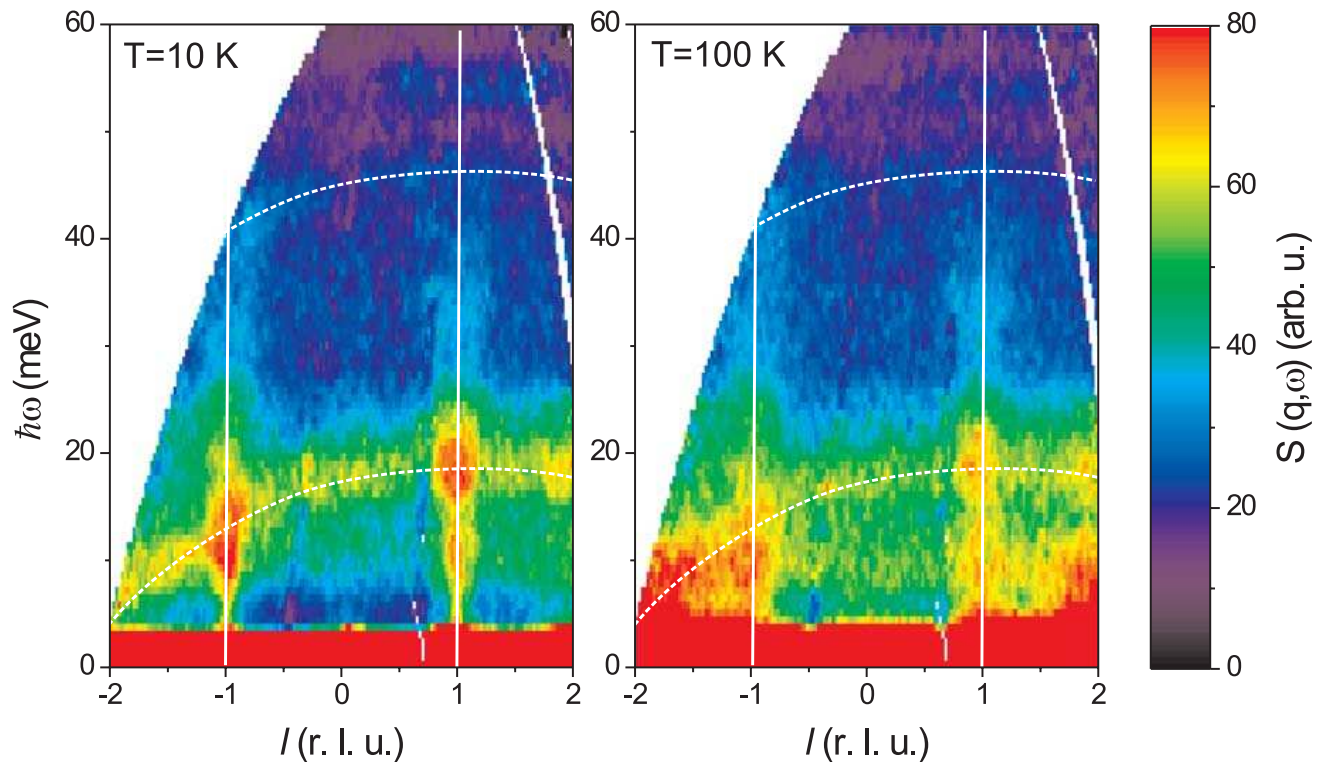


FIG. 3: (Color online) Time of flight spectrum collected in  $\text{BaCu}_2(\text{Si}_{0.5}\text{Ge}_{0.5})_2\text{O}_7$  at  $T = 5$  K using Setup 2, in projection onto the  $c$  (chain) axis. Solid and dashed white lines are 1D antiferromagnetic zone-centers for the two crystallographic domains present in the sample, respectively.

this geometry is optimal for achieving a high wave vector resolution along the chains. A typical constant- $E$  scan of this type measured using Setup 1 at  $\hbar\omega = 12$  meV is shown in Fig. 1a. An alternative approach is to restrict data collection to  $\mathbf{q} = (h \pm x, k, l \pm x)$ , where  $x$  is arbitrary and  $h$  and  $l$  are integers. At these positions the scattering in both crystallographic domains occurs at wave vectors that are equivalent in the 1D sense. The presence of two domains still has to be carefully taken into account when calculating the spectrometer resolution function. This second approach is the only option of high energy transfers, where in each domain the scattering is spread out over a wide  $q$ -range, and necessarily interferes with that from the other domain. Constant- $E$  scans collected in this mode are shown in Fig. 1b,c. A constant background was subtracted from all 3-axis scans shown.

At low temperatures the magnetic dynamic structure factor of disorder-free quantum  $S = 1/2$  chains can be accurately approximated by the Müller Ansatz<sup>5,13,17,18,19,20,21</sup> (MA). For  $\text{BaCu}_2(\text{Si}_{0.5}\text{Ge}_{0.5})_2\text{O}_7$ , in Ref. 6 it was claimed that in constant- $E$  scans the magnetic scattering appears broader than the MA expectation, and is more consistent with Lorentzian line shapes. It was since realized that the elongated Lorentzian-type “tails” are a result of background contamination.<sup>11</sup> Analyzing the new data we find that the MA peak profiles

actually work for  $\text{BaCu}_2(\text{Si}_{0.5}\text{Ge}_{0.5})_2\text{O}_7$  rather well. For each measured constant- $E$  scan the MA dynamic structure factor was numerically folded with the known spectrometer resolution function and fit to the experimental data. The DCP spin wave velocity  $v$  was fixed at  $\pi\langle J \rangle c/4 = 404$  meVÅ. The only variable parameter was an overall scaling factor. Excellent fits were obtained at all energies (solid lines and shaded areas in Fig. 1). Simulations using Lorentzians convoluted with the experimental resolution function are shown in dashed lines. Lorentzians provide satisfactory fits only at low energy transfers, where the finer spectral features are masked by the limited experimental resolution anyway. At high energies deviations from Lorentzian line shapes become obvious, and a flattening at the intensity maximum resulting in typical “top-hat” shapes is clearly observed (Fig. 1c).

## B. TOF data

Specifics of the TOF geometry make separating the signals originating from the two crystallographic domains less straightforward. Typical data collected using Setups 2 and 3 are shown in Figs. 2 and 3 in projection onto the  $c$ -axis of one of the domains (domain A). Vertical streaks of scattering at the 1D AF zone-centers  $l = \pm 1$  for

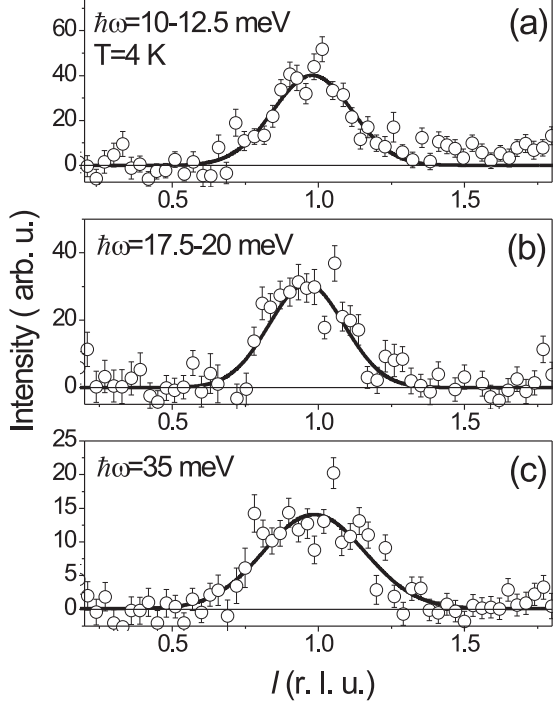


FIG. 4: Typical constant- $E$  cuts of the TOF data for  $\text{BaCu}_2(\text{Si}_{0.5}\text{Ge}_{0.5})_2\text{O}_7$  collected using Setup 2 at  $T = 4$  K. Solid lines are Gaussian fits. A flat background has been subtracted.

this domain are clearly visible and highlighted by vertical solid lines. However, they are intersected by the 1D AF zone-centers of the 2nd domain (domain B), represented by the dashed curves. The corresponding magnetic scattering from domain B appears as a series of arcs. In order to separated it from domain-A scattering one has to rely on certain assumptions and approximations. At small energy transfers, where all the scattering is concentrated in the vicinity of the 1D zone-centers, one can simply regard all data collected sufficiently far from domain B 1D zone-centers as belonging to domain A. Constant-energy cuts obtained from Setup 2 data using this assumption are shown in Fig. 4 (constant background subtracted). Compared to 3-axis data, the experimental wave vector resolution of the TOF setup is insufficient to resolve any fine structure in the scattering profile.

An alternative approach is to select a geometry in which domain B 1D zone-centers cross that of domain A at an almost constant energy, as in the case of  $l = 1$  in Fig. 3. For a constant- $E$  cut extracted from such a data set, the magnetic scattering by domain B will, to a good approximation, correspond to a constant  $l$ -value. Since 1D magnetic scattering is independent of  $h$  and  $k$ , it will amount to a constant background contribution. Constant-energy scans extracted from Setup 3 data us-

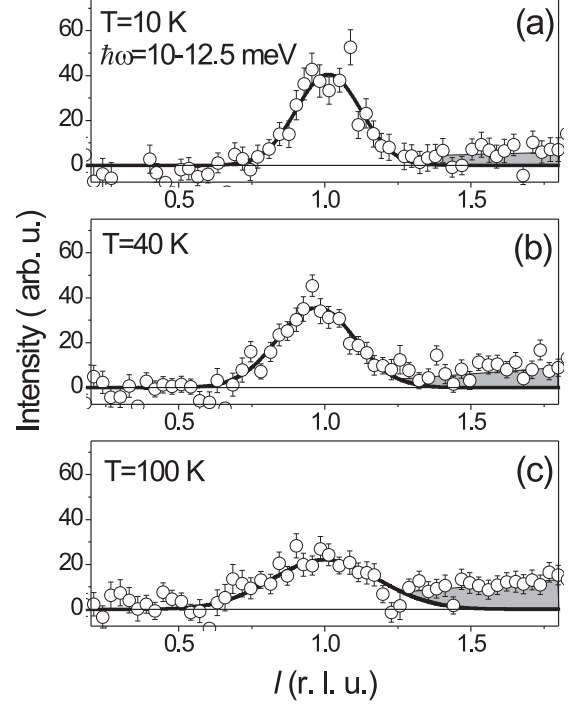


FIG. 5: Temperature evolution of a constant- $E$  cut measured for  $\text{BaCu}_2(\text{Si}_{0.5}\text{Ge}_{0.5})_2\text{O}_7$  using Setup 3. Solid lines are Gaussian fits. A flat background has been subtracted. The shaded area is due to scattering by phonons.

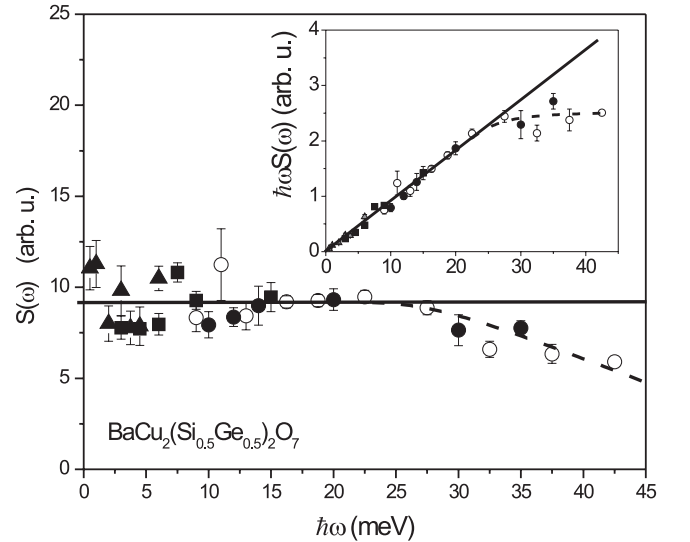


FIG. 6: Scaling of  $S(\omega)$  measured in  $\text{BaCu}_2(\text{Si}_{0.5}\text{Ge}_{0.5})_2\text{O}_7$  (symbols). Solid circles: Setup 1 (3-axis),  $T = 12$  K. Open circles: setup 2 (TOF),  $T = 5.5$  K. Squares: Ref.6, re-analyzed thermal-neutron 3-axis data,  $T = 12$  K. Triangles: Ref.6, cold-neutron 3-axis data,  $T = 1.5$  K. Solid line: Luttinger spin liquid at  $T \rightarrow 0$ . The dashed curves are guides for the eye. Inset: same plot for  $\hbar\omega S(\omega)$ .

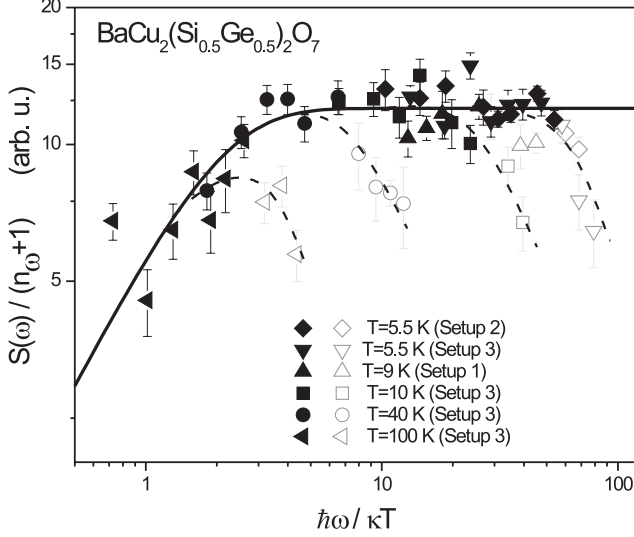


FIG. 7: Measured temperature scaling of  $S(\omega)/(n_\omega + 1)$  for  $\text{BaCu}_2(\text{Si}_{0.5}\text{Ge}_{0.5})_2\text{O}_7$ . Open symbols correspond to energy transfers above 25 meV. The solid line is the approximate scaling function for a Luttinger spin liquid.<sup>17</sup> Dashed lines are guides for the eye and emphasize the breakdown of scaling at high energies.

ing this procedure are shown in Fig. 5. A temperature-induced broadening of the scattering profiles is clearly visible. Data analysis at high temperatures is complicated by the increasing phonon contribution to the background (Fig. 3b and shaded areas in Fig. 5). For this reason, the measurements were limited to  $T \lesssim 100$  K.

### C. Scaling quantities

The central purpose of this study was investigating the energy and temperature scaling of the spin correlations in  $\text{BaCu}_2(\text{Si}_{0.5}\text{Ge}_{0.5})_2\text{O}_7$ . Much of the existing experimental work concentrated on the scaling of the dynamic structure measured at the 1D zone-center  $S(\pi, \omega)$ .<sup>12,13</sup> This quantity is defined exclusively by long wavelength AF spin correlations. In this regime quantum spin chains are equivalent to a Luttinger spin liquid. The existing exact analytical predictions for this model,<sup>22</sup> make a comparison with theory straightforward. The problem, however, is that measuring  $S(\pi, \omega)$  requires wave vector selectivity: the experimental wave vector resolution  $\delta q$  must be small compared to the  $q$ -width of scattering at a particular energy transfer. The latter can be estimated as  $\hbar\omega/v$ , where  $v$  is the DCP spin wave velocity. Thus, there will always be a lower limit on energy transfer for the measurements, determined by the experimental resolution:  $\hbar\omega_{\min} \sim \delta q v$ . For our 3-axis setup, the practical lower limit is about 20 meV, and is still higher for the TOF experiments. A more robust approach is to measure the  $q$ -integrated structure factor  $S_0(\omega) = \int S(q, \omega) d\omega$ .<sup>12</sup> In the hydrodynamic limit the integral is to be taken

over the entire  $q$ -range, but, on a lattice, it should be limited to the vicinity of the 1D AF zone-center. Taking the integral entirely eliminates the effect of wave vector resolution. In our case of  $\text{BaCu}_2(\text{Si}_{0.5}\text{Ge}_{0.5})_2\text{O}_7$ ,  $S(\omega)$  was determined from the cross section fits to individual 3-axis scans  $q$ -scans (solid lines in Fig. 1). For the TOF data we employed empirical Gaussian fits to constant- $E$  cuts, as shown in solid lines in Figs. 4 and 5. The bulk of the low-temperature data for  $S(\omega)$  are shown in Fig. 6 (circles). Re-analyzed data from Ref. 6 are plotted as solid squares (thermal neutron measurements) and triangles (cold neutrons). The inset shows the same data scaled by the energy transfer. All  $S(\omega)$  data collected in this work at different temperatures, after an appropriate scaling (see below), are plotted in Fig. 7. Solid symbols are for energy transfers  $\hbar\omega < 25$  meV, while open symbols represent higher energies.

## IV. DISCUSSION

The finite-temperature scaling for the dynamic structure factor of a Luttinger spin liquid is given by:<sup>22</sup>

$$S(\tilde{q}, \omega) \propto (n_\omega + 1) \frac{1}{T} F\left(\frac{\hbar\omega}{\kappa T}, \frac{v\tilde{q}}{\kappa T}\right), \quad (1)$$

$$F(x, y) = \text{Im} \left( \frac{\Gamma(\frac{1}{4} - i\frac{x-y}{4\pi}) \Gamma(\frac{1}{4} - i\frac{x+y}{4\pi})}{\Gamma(\frac{3}{4} - i\frac{x-y}{4\pi}) \Gamma(\frac{3}{4} - i\frac{x+y}{4\pi})} \right), \quad (2)$$

where  $(n_\omega + 1) = (1 - e^{-\hbar\omega/(\kappa T)})^{-1}$ . For an  $S = 1/2$  Heisenberg spin chain, this scaling applies to AF spin correlations near a 1D AF zone-center  $q = q_0$ , so that  $\tilde{q} = q - q_0$ . The integral over  $\tilde{q}$  needed to derive the scaling for  $S(\omega)$  cannot be easily taken analytically. However, it can be very accurately approximated by:<sup>17</sup>

$$S(\omega) \sim \tanh\left(\frac{\hbar\omega}{2\kappa T}\right). \quad (3)$$

To a very good approximation, then,  $S(\omega) = \text{const}$  at  $\hbar\omega \gg \kappa T$ . The same result can be obtained by integrating the Muller Ansatz, which at small  $\tilde{q}$  and  $\hbar\omega \ll \hbar\omega_{\text{ZB}}$  coincides with the  $T \rightarrow 0$  asymptotic form for the scaling function 2.

The low-temperature data plotted in Fig. 6 show that in  $\text{BaCu}_2(\text{Si}_{0.5}\text{Ge}_{0.5})_2\text{O}_7$ ,  $S(\omega)$  is indeed practically energy independent up to about 25 meV energy transfer (solid lines), *i. e.*, up to about half of the zone-boundary energy  $\hbar\omega_{\text{ZB}}$ . In this energy range the measured temperature dependence of  $S(\omega)$  is consistent with the scaling form 3, as shown in Fig. 7, solid line. This plot, with a dynamic range for  $\hbar\omega/\kappa T$  from 1 to 100, is complimentary to that shown in Ref. 17 for Cu-benzoate with  $\hbar\omega/\kappa T$  in the range of about 0.1 to 10. Above 25 meV, the measured  $S(\omega)$  starts to decrease and  $\omega S(\omega)$  deviates from linear behavior, even at low temperatures (Fig. 6, dashed lines). This breakdown of scaling is observed at all temperatures, as shown by the open symbols and dashed lines in Fig. 7.



The observed breakdown of scaling for  $S(\omega)$  at about half the zone-boundary energy is rather interesting, if not unexpected. Work on  $\text{KCuF}_3$  has shown that for  $S(\pi, \omega)$  the scaling relation 1 holds at much higher energies, up to about 140% of  $\hbar\omega_{\text{ZB}}$ .<sup>13</sup> Of course, the difference is due to the fact that Eq. 1 is derived for the continuum limit  $|\tilde{q}| \ll 1$ . Unlike  $S(\pi, \omega)$ ,  $S(\omega)$  includes scattering at wave vectors  $-\hbar\omega/v < \tilde{q} < \hbar\omega/v$ , and thus rapidly exceeds the domain of validity of Eq. 1 as  $\hbar\omega/v$  increases. Note that the existing  $S(\omega)$  scaling data for Cu-benzoate also extend to about half of  $\hbar\omega_{\text{ZB}}$ .

## V. CONCLUDING REMARKS

It is now clear that at energy transfers exceeding about 0.5 meV,  $\text{BaCu}_2(\text{Si}_{0.5}\text{Ge}_{0.5})_2\text{O}_7$  behaves as a prototypical disorder-free 1D quantum  $S = 1/2$  system. In this sense, the claim of Ref. 7 that  $\text{BaCu}_2(\text{Si}_{1-x}\text{Ge}_x)_2\text{O}_7$  com-

pounds are useful  $S = 1/2$  chain materials, with a continuously adjustable exchange constant, is fully justified. On the other hand, the suppression of the ordering temperature, the anomalous low-temperature increase in the bulk susceptibility,<sup>6</sup> recent ESR data<sup>23</sup>, and preliminary neutron spin echo measurements<sup>24</sup> clearly show that, at lower energies (longer time scales), quenched disorder *does* become relevant. Understanding the scaling of spin correlations in this regime is a challenge for future studies.

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